EXHIBIT 146

To: Koumou, Janis[Koumou.Janis@Endo.com]; Connell, Jill[Connell.Jill@Endo.com]
From: Walker, Lisa
Sent: Tue 3/20/2012 12:18:52 AM
Subject: FW: Annual Training Curriculum Review and Approval for Required Training
Customer Service 2012 Curriculum.xlsx
Master Effective Procedural Documents as of 2-8-2012.xlsx

E0674.1

No changes are required, I will sign and scan to you tomorrow.

Thanks,

Lisa

From: Koumou, Janis
Sent: Monday, March 19, 2012 11:23 AM
To: Connell, Jill; Walker, Lisa
Subject: FW: Annual Training Curriculum Review and Approval for Required Training

Hi Jill and Lisa,

Just another reminder. These should be finalized by the end of this month.

Thanks,

Janis D. Koumou
516-247-5701 x5735
Koumou.janis@endo.com

endo Pharmaceuticals

From: Koumou, Janis
Sent: Tuesday, March 06, 2012 2:35 PM
To: Connell, Jill; Walker, Lisa
Subject: FW: Annual Training Curriculum Review and Approval for Required Training

Reminder – Due Date 3/15/2012

Janis D. Koumou
516-247-5701 x5735
Koumou.janis@endo.com

endo Pharmaceuticals

From: Koumou, Janis
Sent: Thursday, February 09, 2012 8:35 AM
To: Connell, Jill; Walker, Lisa
Subject: Annual Training Curriculum Review and Approval for Required Training

Hi Jill and Lisa,

It is time to conduct the annual review of the Training Curricula for your employees. An Excel spreadsheet is attached with ***separate worksheets for each job title*** within your functional area.
Please review each curricula for additions or deletions of procedures as you feel are ***appropriate and required for each job function***. A Master List of current Procedural Documents (as of 2/8/2012) is included for your convenience, however you may want to check the EDGAR system to ensure that you have identified all procedures for required training. Mandatory training is noted in green highlighting.

Actions Required By You:
If no changes are required, please print, sign, and date each worksheet and send the ***hard copy signatures*** back to me via interoffice mail to the address indicated below. E06742
If changes are required, please **clearly** indicate what you wish added or removed and return the file back to me via email. I will update them accordingly and send you a new copy for your signature.
Due Date: March 15, 2012

Background:
To paraphrase 21 CFR 211.25 Personnel Qualifications, each person must have sufficient education, experience, and training, or a combination of these, to perform their job functions. To meet these criteria, training curricula have been developed for each job title/function so that all employees are periodically trained on those procedures that govern their specific job functions and have the knowledge required to be qualified to execute the tasks involved. A curriculum also ensures that employees receive training on revisions made to these procedures during the year (excluding grammatical, formatting, etc.) so that they remain in compliance with the current process.

If you have any questions, please do not hesitate to ask.

Regards,

Janis Koumou
QA Compliance Specialist
865 Merrick Ave.
Westbury, NY 11590
Koumou.janis@endo.com
516-247-5735
Fax 516-247-5748

2012 Annual Curriculum Review

Customer Service - Associate Director

Authorized Distributor of Record	Tech-Ops-SOP-00004
Internal Procedural Document Processing Process	QA-SOP-00057
Lidoderm Lifetime Supply Program	MI-BP-903
Processing Internal Procedural Documents Using EDGAR	QA-SOP-WI-00003
Vantas Lifetime Supply Program	MI-BP-904

Mandatory training for all on-site employees.	
Adverse Event Reporting	EndoAE01
Procedure for Food and Drug administration (FDA) Inspections	QA-SOP-00012
Record Management Policy	Legal-Policy-00007

Approved By Date

2012 Annual Curriculum Review

Customer Service - Specialist

Authorized Distributor of Record	Tech-Ops-SOP-00004
Lidoderm Lifetime Supply Program	MI-BP-903
Vantas Lifetime Supply Program	MI-BP-904

Mandatory training for all on-site employees.	
Adverse Event Reporting	EndoAE01
Procedure for Food and Drug administration (FDA) Inspections	QA-SOP-00012
Record Management Policy	Legal-Policy-00007

Approved By Date

E0674.5

Produced In Native Format

object_name	title	department	r_version_label	a_effective_date
IM-SOP-WI-00004	Electronic Systems General Reporting	Information Management	Effective, CURRENT, 1.0	11/3/2009 16:10
IM-SOP-WI-00005	Conducting a Root Cause Analysis	Information Management	Effective, CURRENT, 1.0	11/3/2009 16:10
IM-SOP-WI-00006	SAP Problem Reporting	Information Management	Effective, CURRENT, 1.0	11/3/2009 16:10
IM-SOP-WI-00007	CSV Test Script Execution and Documentation	Information Management	CURRENT, Effective, 2.0	11/22/2011 10:04
IM-SOP-WI-00008	IM GxP Compliance Assessments	Information Management	CURRENT, Effective, 2.0	7/21/2010 11:51
IM-SOP-WI-00009	Create Validation Deliverables	Information Management	CURRENT, Effective, 2.0	11/22/2011 10:06
IM-SOP-WI-00011	Window Server Patch Management	Information Management	Effective, CURRENT, 1.0	2/4/2010 15:55
IM-SOP-WI-00014	Changes to GxP Validated Systems	Information Management	CURRENT, Effective, 2.0	10/11/2010 7:42
Med-BP-WI-00006	Development of Recruitment and Retention Plan-In House	Clinical	Effective, CURRENT, 1.0	7/13/2009 11:15
Nonclinical-SOP-WI-00001	Considerations for Impurity Safety Evaluation in Clinical Trials Material	Nonclinical	Effective, CURRENT, 1.0	1/13/2011 9:59
PD-BP-WI-00001	Manufacturing, Analytical, and Clinical Supplies Processes in Pharmaceutical Development	Pharmaceutical Development	Effective, CURRENT, 1.0	8/12/2010 16:01
PD-BP-WI-00002	Drug Development Process	Pharmaceutical Development	Effective, CURRENT, 1.0	8/12/2010 16:01
PD-BP-WI-00004	Sourcing of Materials	Pharmaceutical Development	Effective, CURRENT, 1.0	8/12/2010 16:01
QA-SOP-WI-00001	Processing External Documents Generated by Endo Contractors through EDGAR	Quality Assurance	CURRENT, Effective, 8.0	11/8/2011 10:17
QA-SOP-WI-00003	Processing Internal Procedural Documents Using EDGAR	Quality Assurance	CURRENT, Effective, 11.0	10/14/2011 9:18
QA-SOP-WI-00004	Processing Internal CMC Documents in EDGAR	Quality Assurance	CURRENT, Effective, 7.0	11/8/2011 10:17
QA-SOP-WI-00005	Entering Approved Documents into EDGAR	Quality Assurance	CURRENT, Effective, 11.0	8/9/2011 14:35
QA-SOP-WI-00006	Product Disposition Work Instructions	Quality Assurance	CURRENT, Effective, 7.0	9/21/2011 14:42
QA-SOP-WI-00009	Labeling Control Work Instructions	Quality Assurance	CURRENT, Effective, 3.0	10/27/2010 12:04
QA-SOP-WI-00010	Processing External Documents Generated by Endo Contractors in the Manual System	Quality Assurance	CURRENT, Effective, 4.0	4/19/2011 16:29
QA-SOP-WI-00011	Processing Product Complaints	Quality Assurance	CURRENT, Effective, 8.0	7/7/2011 9:08
QA-SOP-WI-00013	Product Inspection and Release from DC	Quality Assurance	CURRENT, Effective, 4.0	10/27/2010 12:04
QA-SOP-WI-00014	ComplianceWire Work Instructions for Adminsitrators	Quality Assurance	CURRENT, Effective, 2.0	8/15/2011 13:41
RA-BP-WI-00004	WI - Brief Summaries	Regulatory Affairs	Effective, CURRENT, 1.0	9/15/2008 11:31

object_name	title	department	r_version_label	a_effective_date
IM-SOP-Form-00001	Electronic System Evaluation Checklist For 21 CFR Part 211 Compliance	Information Management	CURRENT, Effective, 2.0	11/20/2008 17:09
IM-SOP-Form-00002	Electronic System Evaluation Checklist For 21 CFR Part 11 Compliance	Information Management	CURRENT, Effective, 2.0	11/20/2008 17:09
IM-SOP-Form-00004	Electronic System Request Form	Information Management	CURRENT, Effective, 12.0	11/7/2011 9:25
IM-SOP-Form-00006	Electronic Systems User Account Authorization	Information Management	CURRENT, Effective, 4.0	8/2/2010 12:54
IM-SOP-Form-00008	Data Archive Request Form	Information Management	CURRENT, Effective, 2.0	11/20/2008 17:09
IM-SOP-Form-00010	SAP Account Authorization Form	Information Management	CURRENT, Effective, 2.0	5/8/2008 9:01
IM-SOP-Form-00011	SAP Change Request Form	Information Management	CURRENT, Effective, 2.0	5/8/2008 9:01
IM-SOP-Form-00012	SAP Transport Request Form	Information Management	CURRENT, Effective, 3.0	3/22/2011 13:35
IM-SOP-Form-00013	Endo SAP Job Scheduling Request	Information Management	CURRENT, Effective, 4.0	12/21/2010 9:36
IM-SOP-Form-00016	IM Vendor Qualification Form	Information Management	Effective, CURRENT, 1.0	5/28/2009 10:58
IM-SOP-Form-00020	Form-fishbone template	Information Management	Effective, CURRENT, 1.0	11/3/2009 16:10
IM-SOP-Form-00021	Endo GxP Electronic Systems List	Information Management	CURRENT, Effective, 5.0	7/15/2011 11:51
IM-SOP-Form-00022	Compliance Assessment Form	Information Management	CURRENT, Effective, 7.0	11/7/2011 8:49
IM-SOP-Form-00023	Validation Deliverables Worksheet	Information Management	CURRENT, Effective, 5.0	11/7/2011 8:48
Med-BP-Form-00001	Presentation Request Form	Clinical	Effective, CURRENT, 1.0	5/19/2008 17:32
Med-BP-Form-00002	Presentation Request Form Checklist	Clinical	Effective, CURRENT, 1.0	5/19/2008 17:32
Med-BP-Form-00006	PVRM Notification of Premature Unblinding	Clinical	Effective, CURRENT, 1.0	4/27/2009 18:32
Med-BP-Form-00011	Vantas Lifetime Supply - Confirmation Letter Template	Clinical	Effective, CURRENT, 1.0	10/29/2009 13:52
Med-SOP-Form-00044	Phase I Monitoring Plan Template	Clinical	Effective, CURRENT, 1.0	10/24/2008 15:56
PD-SOP-Form-00001	Equipment Calibration Form	Generic Development	CURRENT, Effective, 2.0	4/5/2010 13:17
PD-SOP-Form-00003	Analytical Request Form	Generic Development	CURRENT, Effective, 3.0	8/4/2010 11:19
PD-SOP-Form-00005	Checklist For Laboratory Related Errors	Generic Development	CURRENT, Effective, 2.0	6/16/2010 15:54
PD-SOP-Form-00006	Peripheral Equipment Performance Verification Form	Generic Development	CURRENT, Effective, 2.0	4/5/2010 13:24
PD-SOP-Form-00007	Balance Daily Weight Check Form	Generic Development	CURRENT, Effective, 2.0	4/5/2010 13:18
PD-SOP-Form-00008	pH Meter Daily Log Book Form	Pharmaceutical Development	Effective, CURRENT, 1.0	11/6/2006 0:00
PD-SOP-Form-00009	Performance Verification/Calibration Form	Generic Development	CURRENT, Effective, 2.0	4/5/2010 13:20
PD-SOP-Form-00010	Controlled Substance Laboratory Use Record	Pharmaceutical Development	Effective, CURRENT, 1.0	7/2/1998 0:00
PD-SOP-Form-00011	Employee Acknowledgment of Understanding and Compliance-Controlled Substances	Pharmaceutical Development	Effective, CURRENT, 1.0	3/22/2004 0:00
PD-SOP-Form-00012	Request for ECSTA Number	Pharmaceutical Development	CURRENT, Effective, 2.0	6/20/2009 16:35
PD-SOP-Form-00013	Controlled Substance Transfer Voucher	Pharmaceutical Development	Effective, CURRENT, 1.0	7/12/2004 0:00
PD-SOP-Form-00014	Transfer of License Form for CII-V Material Only	Pharmaceutical Development	Effective, CURRENT, 1.0	3/23/2004 0:00
PD-SOP-Form-00015	Controlled Substance Laboratory Request Record	Pharmaceutical Development	Effective, CURRENT, 1.0	3/23/2004 0:00
PD-SOP-Form-00016	Controlled Substance Material Use Record	Pharmaceutical Development	Effective, CURRENT, 1.0	3/23/2004 0:00
PD-SOP-Form-00017	Controlled Substance Seal Form	Pharmaceutical Development	Effective, CURRENT, 1.0	10/8/2003 0:00
PD-SOP-Form-00023	Incident Investigation Report Environmental Health and Safety	Pharmaceutical Development	Effective, CURRENT, 1.0	5/10/2000 0:00
PD-SOP-Form-00024	Cancellation of Stability Study Notices	Generic Development	CURRENT, Effective, 3.0	4/5/2010 13:20
PD-SOP-Form-00025	Out of Alert Notice	Generic Development	CURRENT, Effective, 2.0	4/5/2010 13:22
PD-SOP-Form-00026	Stability Sample Information	Generic Development	CURRENT, Effective, 2.0	3/12/2010 7:51
PD-SOP-Form-00028	Stability Study Waiver Form	Pharmaceutical Development	Effective, CURRENT, 1.0	4/3/2007 0:00
PD-SOP-Form-00029	Out of Specification Notice	Generic Development	CURRENT, Effective, 2.0	3/12/2010 7:51
PD-SOP-Form-00030	Stability Study Assessment Form	Pharmaceutical Development	Effective, CURRENT, 1.0	1/24/2002 0:00
PD-SOP-Form-00034	ANALYTICAL DEVELOPMENT ELECTRONIC SYSTEM USER ACCOUNT AUTHORIZATION FORM	Generic Development	Effective, CURRENT, 1.0	7/27/2010 12:38

PD-SOP-Form-00035	GMP Sample Usage Accountability Form	Generic Development	Effective, CURRENT, 1.0	8/26/2010 11:45
PD-SOP-Form-00036	Failed Run Tracking Form	Generic Development	Effective, CURRENT, 1.0	11/18/2010 8:48
PD-SOP-Form-00037	Verification of Compendial Methods - Assessment of Requirements	Generic Development	Effective, CURRENT, 2.0	3/11/2011 14:27
PM-SOP-Form-00015	Quality Complaint	Pharmaceutical Development	CURRENT, Effective, 2.0	5/30/2008 9:07
QA-SOP-Form-00004	Record of Training	Quality Assurance	Effective, CURRENT, 2.0	9/7/2007 8:34
QA-SOP-Form-00006	Products Complaint Summary Form	Quality Assurance	Effective, CURRENT, 1.0	11/7/2003 0:00
QA-SOP-Form-00007	Document Review Notice	Quality Assurance	CURRENT, Effective, 2.0	4/19/2011 16:29
QA-SOP-Form-00009	Official Copy Distribution Record	Quality Assurance	Effective, CURRENT, 1.0	1/16/2001 0:00
QA-SOP-Form-00011	Investigation Report Approval Form	Quality Assurance	CURRENT, Effective, 5.0	7/5/2011 15:44
QA-SOP-Form-00015	Investigation/ Deviation Close-Out Report Form	Quality Assurance	CURRENT, Effective, 3.0	7/5/2011 15:47
QA-SOP-Form-00017	Product Disposition Notification for Physician Samples	Quality Assurance	CURRENT, Effective, 4.0	6/3/2011 8:37
QA-SOP-Form-00018	Annual Product Report Review Form	Quality Assurance	CURRENT, Effective, 2.0	11/12/2009 9:41
QA-SOP-Form-00019	Finished Goods Inspection Report For Tablets and Capsules	Quality Assurance	CURRENT, Effective, 5.0	10/27/2010 12:04
QA-SOP-Form-00021	Internal Product Disposition Notification to Endo Distribution	Quality Assurance	CURRENT, Effective, 3.0	6/3/2011 8:38
QA-SOP-Form-00022	Request for Inspection	Quality Assurance	CURRENT, Effective, 2.0	9/30/2008 9:18
QA-SOP-Form-00023	100% Inspection Report	Quality Assurance	CURRENT, Effective, 2.0	10/15/2008 10:19
QA-SOP-Form-00024	Returned Goods Request for Inspection Form	Quality Assurance	Effective, CURRENT, 1.0	2/19/2007 0:00
QA-SOP-Form-00025	Deviation Report Form	Quality Assurance	CURRENT, Effective, 3.0	7/5/2011 15:50
QA-SOP-Form-00026	QA Batch Record Review Form	Quality Assurance	CURRENT, Effective, 5.0	3/31/2011 10:11
QA-SOP-Form-00028	QA Release Of Clinical Supplies To Clinical Supplies Manager	Quality Assurance	CURRENT, Effective, 3.0	3/31/2011 10:12
QA-SOP-Form-00030	Notice Of Product Reject	Quality Assurance	Effective, CURRENT, 1.0	3/16/2007 0:00
QA-SOP-Form-00032	Approval Form	Quality Assurance	CURRENT, Effective, 4.0	11/8/2011 10:17
QA-SOP-Form-00034	Full Service Employee signature Record	Quality Assurance	CURRENT, Effective, 3.0	3/5/2010 17:14
QA-SOP-Form-00038	Document Obsoletion Checklist	Quality Assurance	CURRENT, Effective, 4.0	10/14/2011 9:18
QA-SOP-Form-00039	Aborted Document Checklist	Quality Assurance	CURRENT, Effective, 3.0	5/11/2010 10:49
QA-SOP-Form-00042	Document Processing Checklist for Internal Documents Processed by Manual System	Quality Assurance	CURRENT, Effective, 4.0	10/1/2010 14:56
QA-SOP-Form-00043	Document Distribution and Reconciliation Notice	Quality Assurance	CURRENT, Effective, 2.0	8/8/2008 13:43
QA-SOP-Form-00052	Products Complaint Contact Notes	Quality Assurance	CURRENT, Effective, 2.0	6/29/2007 15:04
QA-SOP-Form-00053	Release Product with Superseded Labeling	Quality Assurance	Effective, CURRENT, 1.0	6/21/2004 0:00
QA-SOP-Form-00054	Quality Quarantine Form	Quality Assurance	CURRENT, Effective, 2.0	6/8/2009 17:02
QA-SOP-Form-00056	Retain Sample Inspection for Annual Product Review	Quality Assurance	CURRENT, Effective, 2.0	5/15/2009 16:55
QA-SOP-Form-00057	Finished Goods Inspection Report for Topical Products (Patch, Ointment, Cream, and Gel)	Quality Assurance	CURRENT, Effective, 8.0	10/27/2010 12:04
QA-SOP-Form-00058	Finished Goods Inspection Report for Syrup Products	Quality Assurance	CURRENT, Effective, 5.0	10/27/2010 12:04
QA-SOP-Form-00059	Finished Goods Inspection Report for Vial and Ampul Products	Quality Assurance	CURRENT, Effective, 6.0	10/27/2010 12:04
QA-SOP-Form-00061	Packaging Batch Record Review Checklist	Quality Assurance	CURRENT, Effective, 2.0	1/10/2008 16:44
QA-SOP-Form-00062	Manufacturing Batch Record Review Checklist	Quality Assurance	CURRENT, Effective, 2.0	1/10/2008 16:42
QA-SOP-Form-00063	Expiration Date Change Notification	Quality Assurance	Effective, CURRENT, 1.0	11/14/2005 0:00
QA-SOP-Form-00064	Product Release Authorization to Endo Distributor	Quality Assurance	CURRENT, Effective, 3.0	1/6/2010 11:56
QA-SOP-Form-00065	Contractor Screening Questionnaire & Follow Up	Quality Assurance	CURRENT, Effective, 3.0	5/20/2011 13:42
QA-SOP-Form-00066	API Supplier Screening Questionnaire & Follow Up	Quality Assurance	CURRENT, Effective, 2.0	1/29/2010 8:31
QA-SOP-Form-00067	Contract Manufacturer Quality Performance Report Card	Quality Assurance	CURRENT, Effective, 3.0	4/27/2009 17:18
QA-SOP-Form-00070	Independent Contractor Signature Record	Quality Assurance	CURRENT, Effective, 2.0	3/5/2010 17:13
QA-SOP-Form-00071	Stability Sample Information for Clinical Supply Release	Quality Assurance	CURRENT, Effective, 3.0	4/4/2011 16:28

QA-SOP-Form-00074	Endo Quality Compliance Checklist for Product Complaints	Quality Assurance	CURRENT, Effective, 7.0	2/4/2011 10:51
QA-SOP-Form-00077	Label Activation Request Form	Quality Assurance	CURRENT, Effective, 2.0	1/13/2010 13:51
QA-SOP-Form-00078	Quarantine Shipment Form, Clinical Trial Materials	Quality Assurance	Effective, CURRENT, 1.0	7/29/2008 16:56
QA-SOP-Form-00079	USP Update Product Impact Assessment	Quality Assurance	Effective, CURRENT, 1.0	8/7/2009 9:11
QA-SOP-Form-00080	Request for a New, Revision or Addendum Document	Quality Assurance	CURRENT, Effective, 2.0	1/26/2011 14:09
QA-SOP-Form-00081	Request to Abort or Obsolete Document	Quality Assurance	CURRENT, Effective, 2.0	10/14/2011 9:18
QA-SOP-Form-00082	Endo Clinical QA Document Audit/Review Form	Quality Assurance	Effective, CURRENT, 1.0	9/13/2010 10:15
QA-SOP-Form-00083	Finished Goods inspection Report for Product Kit	Quality Assurance	CURRENT, Effective, 2.0	10/27/2010 12:05
QA-SOP-Form-00084	Product Transportation Requirement Form	Quality Assurance	CURRENT, Effective, 2.0	10/13/2010 9:45
QA-SOP-Form-00087	Release Notification for Products from ENDO Facility	Quality Assurance	Effective, CURRENT, 1.0	5/5/2010 16:29
QA-SOP-Form-00088	Expiration Date Assignment	Quality Assurance	Effective, CURRENT, 1.0	7/12/2010 15:50
QA-SOP-Form-00089	Audit Waiver Form	Quality Assurance	Effective, CURRENT, 1.0	1/29/2010 8:31
QA-SOP-Form-00090	Reduced Sampling Notification Form	Quality Assurance	Effective, CURRENT, 1.0	9/27/2010 14:58
QA-SOP-Form-00091	Master List of Product Expiration Indexes	Quality Assurance	CURRENT, Effective, 7.0	2/1/2012 14:38
QA-SOP-Form-00092	Method Transfer Waiver Form	Quality Assurance	Effective, CURRENT, 1.0	11/29/2010 14:18
QA-SOP-Form-00093	Release Notification For Placebo Samples	Quality Assurance	Effective, CURRENT, 1.0	2/17/2011 14:30
QA-SOP-Form-00094	Material Disposition Form	Quality Assurance	Effective, CURRENT, 1.0	3/9/2011 8:53
QA-SOP-Form-00095	QA Release of ANDA Clinical Supplies and Approval of batches for In Vitro Bioequivalence Studies	Quality Assurance	Effective, CURRENT, 1.0	3/30/2011 10:17
QA-SOP-Form-00096	Medical Device Reporting Decision Tree	Quality Assurance	Effective, CURRENT, 1.0	2/4/2011 10:51
QA-SOP-Form-00104	Discontinuation of Stability Study	Quality Assurance	Effective, CURRENT, 1.0	8/3/2011 9:58
RA-SOP-Form-00006	Submissions Sign Out Sheet	Regulatory Affairs	Effective, CURRENT, 1.0	9/25/2000 0:00
RA-SOP-Form-00009	Clinical Study Site Approved by Regulatory Affairs	Regulatory Affairs	Effective, CURRENT, 1.0	2/20/2001 0:00
RA-SOP-Form-00010	Central Files Document Request	Regulatory Affairs	Effective, CURRENT, 1.0	3/20/2001 0:00
RA-SOP-Form-00011	Central Files Document Request of Original Records	Regulatory Affairs	Effective, CURRENT, 1.0	3/20/2001 0:00
RA-SOP-Form-00015	Regulatory Agency Contact Report Form	Regulatory Affairs	Effective, CURRENT, 1.0	9/10/2001 0:00
RA-SOP-Form-00024	Notification to Implement Change	Regulatory Affairs	Effective, CURRENT, 1.0	12/16/2005 0:00

object_name	title	department	r_version_label	a_effective_date
CD-BP-100	Protocol Development and Protocol Review Committee	Research and Development (R&D)	CURRENT, Effective, 2.0	6/21/2011 20:22
CD-SOP-100	Protocol Development, Review and Approval	Research and Development (R&D)	CURRENT, Effective, 2.0	6/21/2011 20:26
Clinical Affairs Manager Guidance Document December 2011	Clinical Affairs Manager Guidance Document December 2011	Research and Development (R&D)	Effective, CURRENT, 1.0	12/13/2011 9:48
Clinical Affairs Manager Guidance Document Jan 2011	Clinical Affairs Manager Guidance Document Jan 2011	Research and Development (R&D)	Effective, CURRENT, 1.0	4/4/2011 9:43
DM-BP-500	Excel to SAS Data Set Conversion	Research and Development (R&D)	Effective, CURRENT, 1.0	4/5/2011 15:28
DM-BP-501	Reformatting of Data	Research and Development (R&D)	Effective, CURRENT, 1.0	4/5/2011 15:29
DM-BP-502	Development of an Output From Raw Data	Research and Development (R&D)	Effective, CURRENT, 1.0	4/5/2011 15:30
Legal-BP-00003	BP for Record Archiving and Retrieval	Legal	CURRENT, Effective, 2.0	1/4/2011 9:39
Legal-BP-00005	BP for Records Cleanout	Legal	Effective, CURRENT, 1.0	1/4/2011 9:40
LM-BP-800	External Scientific Presentations: Preparation and Execution of Solicited, Unsolicited or Non-Customer Presentations	Research and Development (R&D)	Effective, CURRENT, 1.0	12/16/2010 14:02
LM-BP-801	Creation, Review and Approval of External Scientific Presentations in Response to an Unsolicited Request from an External Individual or Organization	Research and Development (R&D)	Effective, CURRENT, 1.0	1/6/2011 13:51
LM-BP-802	Review and Handling of Investigator Initiated Research Requests	Research and Development (R&D)	Effective, CURRENT, 1.0	1/6/2011 13:51
LM-BP-803	Document Data Room	Research and Development (R&D)	Effective, CURRENT, 1.0	2/15/2011 14:06
Med-BP-00006	Acquisition of New Product Safety Data by Endo PVRM	Clinical	CURRENT, Effective, 3.0	9/24/2009 13:26
Med-BP-00010	Development of Publications	Clinical	Effective, CURRENT, 1.0	5/15/2009 8:56
Med-BP-00011	Development of a Clinical Development Plan	Clinical	Effective, CURRENT, 1.0	5/1/2009 11:37
Med-BP-00015	Development of a Recruitment and Retention Plan	Clinical	Effective, CURRENT, 1.0	4/27/2009 18:19
Med-BP-00027	Educational Grants Administration Process	Clinical	Effective, CURRENT, 1.0	8/31/2010 16:52
Med-BP-00044	Processing of Adverse Event and Product Complaint Reports	Clinical	Effective, CURRENT, 1.0	9/21/2009 16:31
Med-BP-00045	Process for Paper Submission of Adverse Event Reports to FDA	Clinical	CURRENT, Effective, 2.0	1/23/2012 9:57
Med-BP-00046	Triage Process for Adverse Event Reports	Clinical	Effective, CURRENT, 1.0	9/21/2009 16:31
Med-BP-00047	Reconciliation Process for Adverse Event Reports	Clinical	Effective, CURRENT, 1.0	9/21/2009 16:32
Med-BP-00049	Processing of Solicited Adverse Event Reports	Clinical	Effective, CURRENT, 1.0	9/21/2009 16:32
Med-BP-00050	Enhanced Follow-up Process for Adverse Event Reports	Clinical	CURRENT, Effective, 2.0	1/15/2010 15:39
Med-BP-00051	Obtaining Follow-up Information on Safety Reports for Endo Marketed Products	Clinical	Effective, CURRENT, 1.0	9/21/2009 16:33
Med-BP-00052	Manufacturer Notification of Non-Serious and Serious Adverse Event Reports for Non-Endo Products	Clinical	Effective, CURRENT, 1.0	9/21/2009 16:33
Med-BP-00053	Submission Process for IND Safety Reports and NDA Alert Reports for Endo Products	Clinical	Effective, CURRENT, 1.0	9/21/2009 16:33
Med-BP-00055	Clinical Trial Registration	Clinical	Effective, CURRENT, 1.0	12/7/2009 11:51
Med-BP-00056	Processing of a Package Insert Complaint	Clinical	CURRENT, Effective, 2.0	1/13/2010 17:39
MI-BP-900	Referring MI or PVRM Matters to Endo Corporate Compliance.doc	Research and Development (R&D)	Effective, CURRENT, 1.0	10/21/2010 15:51
MI-BP-901	Field Sales Requests for Medical Information/AMCP Dossiers	Research and Development (R&D)	Effective, CURRENT, 1.0	8/8/2011 11:06
MI-BP-902	EMI Document Development and Maintenance	Research and Development (R&D)	Effective, CURRENT, 1.0	10/21/2010 15:52
MI-BP-903	Lidoderm Lifetime Supply Program	Research and Development (R&D)	Effective, CURRENT, 1.0	8/11/2011 20:52
MI-BP-904	Vantas Lifetime Supply Program	Research and Development (R&D)	Effective, CURRENT, 1.0	8/8/2011 11:06
MI-BP-905	Fulfilling Unsolicited Requests for AMCP Dossiers	Research and Development (R&D)	Effective, CURRENT, 1.0	8/8/2011 11:06
MI-BP-906	Receipt of and Response to Inquiries Received by R&D Field Personnel	Research and Development (R&D)	Effective, CURRENT, 1.0	8/11/2011 20:53
Nonclinical-BP-00001	Assignment of Nonclinical Study Numbers	Nonclinical	Effective, CURRENT, 1.0	11/30/2007 8:39
PM-BP-00001	Clinical Supplies Planning	Pharmaceutical Development	CURRENT, Effective, 2.0	12/12/2011 16:12
PV-BP-300	Evaluating Safety Information from Literature Searches	Research and Development (R&D)	Effective, CURRENT, 1.0	2/15/2011 14:07

PV-BP-301	Recalls, Market Withdrawal or Stock Recovery Activities Involving Marketed Products	Research and Development (R&D)	Effective, CURRENT, 1.0	1/23/2012 16:34
QA-BP-00001	Logging and Filing of Executed Batch Records in QD Central Files	Quality Assurance	Effective, CURRENT, 1.0	1/4/2010 11:58
QA-BP-00003	Periodic Monitoring of EDGAR for Compliance	Quality Assurance	Effective, CURRENT, 1.0	6/17/2010 8:23
RA-BP-00014	Brief Summaries	Regulatory Affairs	Effective, CURRENT, 1.0	9/15/2008 11:28
RA-BP-600	Development and Distribution of Changes/Revisions for Approved Product Labeling	Research and Development (R&D)	CURRENT, Effective, 2.0	6/10/2011 9:36
RA-BP-601	Review of Labeling Complaints Unrelated to Product Complaints or Adverse Events	Research and Development (R&D)	Effective, CURRENT, 1.0	11/18/2011 9:12

object_name	title	department	r_version_label	a_effective_date
IM-Policy-00002	IM Security	Information Management	Effective, CURRENT, 1.0	2/4/2010 14:25
Legal-Policy-00003	Orphaned Files Policy	Legal	CURRENT, Effective, 3.0	1/11/2011 14:15
Legal-Policy-00005	Hold Order Policy	Legal	CURRENT, Effective, 2.0	8/31/2009 8:05
Legal-Policy-00006	Backup/Recovery Data Retention Policy	Legal	CURRENT, Effective, 3.0	8/31/2009 8:05
Legal-Policy-00007	Record Management Policy	Legal	CURRENT, Effective, 4.0	1/28/2011 11:13
Legal-Policy-00009	Records Retention Schedule	Legal	CURRENT, Effective, 3.0	8/9/2011 15:30
Med-Policy-00001	Educational Grant Policy	Research and Development (R&D)	CURRENT, Effective, 2.0	2/15/2011 14:08
Med-Policy-00004	Credentialing Policy and Procedure	Research and Development (R&D)	CURRENT, Effective, 2.0	8/11/2011 14:04
Med-Policy-00005	Non-Promotional Review Committee Policy	Research and Development (R&D)	Effective, CURRENT, 1.0	12/15/2011 16:33
QA-Policy-00002	Quality Manual	Quality Assurance	CURRENT, Effective, 10.0	10/18/2011 9:37
QA-Policy-00003	Documentation Management Policy	Quality Assurance	CURRENT, Effective, 3.0	2/23/2010 9:34
QA-Policy-00004	Complaint and Adverse Experience Handling and Reporting Policy	Quality Assurance	CURRENT, Effective, 2.0	11/13/2009 14:08
QA-Policy-00005	Compliance Training Policy	Quality Assurance	CURRENT, Effective, 3.0	12/6/2011 8:53
QA-Policy-00006	Audit Program Policy	Quality Assurance	CURRENT, Effective, 2.0	11/11/2011 9:37
QA-Policy-00008	Confirmatory Testing	Quality Assurance	Effective, CURRENT, 1.0	10/10/2011 9:30
QA-Policy-00010	Communication and Information Sharing Process for Quality & Compliance	Quality Assurance	Effective, CURRENT, 1.0	11/2/2011 14:43

object_name	title	department	a_effective_date	r_version_label
QA-Guideline-00004	Document Naming and Handling Conventions	Quality Assurance	11/2/2011 14:44	Effective, CURRENT, 1.0

object_name	title	department	r_version_label	a_effective_date
QA-Index-00001	Expiration Dating List Voltaren Gel	Quality Assurance	CURRENT, Effective, 3.0	6/10/2010 16:17
QA-Index-00004	Expiration Dating List for Endocet/Percocet	Quality Assurance	CURRENT, Effective, 2.0	3/1/2011 13:54
QA-Index-00005	Expiration Dating List for Endodan/Percodan	Quality Assurance	Effective, CURRENT, 1.0	2/1/2008 0:00
QA-Index-00006	Expiration Dating List for Frova	Quality Assurance	Effective, CURRENT, 1.0	1/28/2008 0:00
QA-Index-00009	Expiration Dating List for Lidoderm	Quality Assurance	CURRENT, Effective, 2.0	6/10/2010 16:17
QA-Index-00010	Expiration Dating List for Moban	Quality Assurance	CURRENT, Effective, 2.0	8/23/2011 14:41
QA-Index-00011	Expiration Dating List for Morphine Sulfate ER	Quality Assurance	Effective, CURRENT, 1.0	9/20/2007 0:00
QA-Index-00014	Expiration Dating List for Zydone Tablets	Quality Assurance	CURRENT, Effective, 2.0	3/1/2011 13:54
QA-Index-00016	Expiration Dating List for Opana Tablets	Quality Assurance	CURRENT, Effective, 2.0	5/15/2009 10:32
QA-Index-00017	Expiration Dating List for Opana ER Tablets	Quality Assurance	CURRENT, Effective, 3.0	6/11/2010 10:06
QA-Index-00018	Expiration Dating List for Opana Injection	Quality Assurance	Effective, CURRENT, 1.0	5/8/2007 0:00
QA-Index-00019	Expiration Dating List Valstar	Quality Assurance	CURRENT, Effective, 2.0	5/26/2011 13:39
QA-Index-00020	Expiration Dating List for Delatestryl	Quality Assurance	Effective, CURRENT, 1.0	9/3/2009 10:08
QA-Index-00023	Expiration Dating List Vantas	Quality Assurance	Effective, CURRENT, 1.0	9/3/2009 10:08
QA-Index-00024	Expiration Dating List for Supprelin LA	Quality Assurance	Effective, CURRENT, 1.0	9/3/2009 10:08
QA-Index-00025	Michophenolate Mofetil Capsules	Quality Assurance	Effective, CURRENT, 1.0	1/22/2010 15:52
QA-Index-00026	Mycophenolate Tablets	Quality Assurance	Effective, CURRENT, 1.0	7/19/2010 11:17
QA-Index-00027	Supppprelin LA Kit	Quality Assurance	Effective, CURRENT, 1.0	6/15/2010 10:20
QA-Index-00028	Expiration dating List for Valstar Kit	Quality Assurance	Effective, CURRENT, 1.0	6/11/2010 10:48
QA-Index-00029	Expiration dating List for Vantas Kit	Quality Assurance	CURRENT, Effective, 2.0	8/31/2011 8:14
QA-Index-00030	Expiration Dating List Losartan Potassium Tablets	Quality Assurance	Effective, CURRENT, 1.0	10/26/2010 14:10
QA-Index-00031	Expiration Dating List Pramipexole Tablets	Quality Assurance	Effective, CURRENT, 1.0	10/27/2010 9:21
QA-Index-00033	Expiration Dating List Fortesta	Quality Assurance	Effective, CURRENT, 1.0	1/4/2011 10:36
QA-Index-00035	Material Index	Quality Assurance	Effective, CURRENT, 1.0	3/9/2011 8:53
QA-Index-00036	Expiration Dating List for Letrozole	Quality Assurance	Effective, CURRENT, 1.0	6/3/2011 11:46
QA-Index-00037	Expiration Dating List for Previfem	Quality Assurance	Effective, CURRENT, 1.0	8/12/2011 16:48
QA-Index-00038	Expiration Dating List for Gildess FE	Quality Assurance	Effective, CURRENT, 1.0	8/22/2011 16:25
QA-Index-00039	Expiration Dating List for Triprevifem	Quality Assurance	Effective, CURRENT, 1.0	8/22/2011 16:25
QA-Index-00040	Expiration Dating List for Cyclafem	Quality Assurance	Effective, CURRENT, 1.0	8/22/2011 16:25

QA-Index-00041	Expiration Dating List for Emoquette	Quality Assurance	Effective, CURRENT, 1.0	8/22/2011 16:25
QA-Index-00042	Expiration Dating List for Orsythia	Quality Assurance	Effective, CURRENT, 1.0	8/22/2011 16:25
QA-Index-00043	Expiration Dating List for Methylprednisolone	Quality Assurance	Effective, CURRENT, 1.0	9/20/2011 11:07
QA-Index-00044	Expiration Dating List for Prednisone	Quality Assurance	Effective, CURRENT, 1.0	9/20/2011 11:07
QA-Index-00045	Expiration Dating List Felodipine ER	Quality Assurance	Effective, CURRENT, 1.0	1/17/2012 13:49